leukemia, athlete's foot, varicose veins, tetanus, typhoid, gonorrhea, staphylococcus, pneumonia, streptothrix, streptococcus, TB virus, carcinoma, sarcoma, treponema, abscess, fistula, hemorrhoids, hernia, irritations, arthritis, bursitis, palsy, diseased lymph nodes, acne, cystitis, boils, bubonic plague, diphtheria, elephantiasis, fungus, impetigo, hardening of the arteries, leprosy, moles, multiple sclerosis, poison oak, poison ivy, poliomyelitis, skin eruptions, spinal meningitis, warts, constipation, typhoid fever, colitis, cataract, glaucoma, leakage of the heart, coronary thrombosis, tetanus, peptic ulcers, and other abnormal and disease conditions.

DISPOSITION: 5–29–61. Default—delivered to the Food and Drug Administration.

6617. Ortho-Structurometer device. (F.D.C. No. 44581. S. No. 44–460 R.)

QUANTITY: One device at Portland, Oreg.

SHIPPED: 4–28–59, from Monrovia, Calif., by Custom Bearings, for J. & E. Enterprises, Inc.

LABEL IN PART: "J. & E. Enterprises, Inc., Model No. Ortho 7 * * * Pasadena, Calif."

ACCOMPANYING LABELING: Leaflets entitled "Self Appraisal" and a posture chart bearing the name "T. E. Hall."

RESULTS OF INVESTIGATION: Examination indicated that the device was a portable unit consisting of two tilt platforms and a control panel for adjusting the platforms to varying degrees. The user stood on the platforms for the intended purpose of changing posture and thereby alleviating various disease and abnormal conditions.

LIBELED: 5–19–60, Dist. Oreg.

CHARGE: 502(a)—when shipped, the labeling of the article contained false and misleading representations that the article was an adequate and effective treatment for preventing or overcoming prolapsed diaphragm, weakened perineum, rectal prolapse, constipation, hemorrhoids, pudendal hemorrhage, prolapsed uterus, hernia, congested uterus and ovaries, tuberculosis, asthma, heart conditions, bladder irritation, femoral hernia, inguinal hernia, visceral ptosis, broken arches, and ear, eye, nose and throat infections, and that the use of the device would correct improper body mechanics and body imbalance to prevent and overcome most common diseases.

DISPOSITION: On 8–30–60, J. & E. Enterprises, Inc., appeared and filed a claim to the device and, on 8–31–60, the cause was removed to the United States District Court for the Northern District of California. On 9–28–60, the claimant filed an answer denying the misbranding. On 5–1–61, the claim and answer were withdrawn and, on 5–25–61, a default decree of forfeiture was filed and the court ordered the device delivered to the Food and Drug Administration.

6618. Vibra-Finger Gum Massager. (F.D.C. No. 45476. S. No. 26–965 R.)

QUANTITY: 31 individually cartoned devices at Los Angeles, Calif., in possession of Gem Products.

SHIPPED: 1–18–61, from New York, N.Y., by Vibra Research Laboratories.

LABEL IN PART: (Ctn.) "Vibra-Finger Professional Gum Massager * * * Distributors Vibra Research Laboratories."

ACCOMPANYING LABELING: Folder in carton reading in part "Your Vibra Finger Gum Massager Instructions For . . ." and leaflets entitled "Vibra-Finger."

RESULTS OF INVESTIGATION: Examination showed that the article was an electrically operated vibrator device, about the size and general shape of an electric razor. A plastic finger-shaped attachment extended from one end of the motor housing. In use, the vibrating plastic finger attachment was pressed against the gums of the mouth.

The above-mentioned leaflets were printed locally at the request of the dealer and were distributed with the device.

LIBELED: 2–16–61, S. Dist. Calif.

CHARGE: 502(a)—when shipped and while held for sale, the accompanying labeling of the article contained false and misleading representations that the article was an adequate and effective treatment for promoting strong healthy gums, preventing loosening of teeth, pyorrhea, and soft irritated gums.

DISPOSITION: 4–17–61. Default—destruction.

6619. Health-Aire device. (F.D.C. No. 44721. S. No. 21–425 R.)

QUANTITY: 20 individually cartoned devices at Akron, Ohio.

SHIPPED: 2–15–60, from Long Island City, N.Y., by Samson United of New York.

LABEL IN PART: (Ctn.) "Health Aire for Mountain Air Purity * * * Samson United of New York F 5966."

ACCOMPANYING LABELING: Instruction sheet in carton reading in part "Instruction Sheet For Your Health-Aire" and folder entitled "Samson's Health-Aire."

RESULTS OF INVESTIGATION: The article was a portable cabinet enclosing a fan, a nylon filter, and an ultraviolet lamp. Room air was circulated through the unit which operated on ordinary household electric current.

LIBELED: 7–14–60, N. Dist. Ohio.

CHARGE: 502(a)—when shipped and while held for sale, the labeling of the article contained false and misleading representations that the article was an adequate and effective treatment for relieving breathing distress and discomfort of allergies, hay fever, asthma, sinus, and colds, and that the article provided health-giving air of "mountain air purity."

DISPOSITION: Samson United Corp. of New York, claimant, having consented to the entry of a decree, judgment of condemnation was entered on 10–14–60, and the court ordered the devices released under bond for relabeling. On 5–2–61, the claimant having failed to file the required bond, the devices were delivered to the Food and Drug Administration.

DRUG FOR VETERINARY USE*

6620. L-K tablets. (F.D.C. No. 45320. S. No. 34–083 R.)

QUANTITY: 6 cases, 12 ctnd. btls. each, at Brooklyn, N.Y.

SHIPPED: During the fall of 1958, from Webster, Mass., by Dr. A. C. Daniels, Inc.

LABEL IN PART: (Ctn.) "Dr. A. C. Daniels L–K Tablets Active Ingredients: Boric Acid, Extract Corn Silk, Extract Hydranges, Extract Buchu, Extract Triticum, Potassium Bicarbonate and Atropine Sulfate 1/2000 grain. Dog–Cat Contents 48 * * * Distributed by Dr. A. C. Daniels, Inc., Boston, Mass. * * * Diuretic. Stimulates the flow of urine. Directions on bottle label."

*See also Nos. 6594, 6602.